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(54) **Aqueous anti-curl fixing fluids**

Wässrige antiroll Fixierflüssigkeit

Liquide de fixation anti enroulement aqueuse

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Description

FIELD OF THE INVENTION

[0001] The present invention relates to cosolvents used in fixers, the cosolvents imparting anti-curl qualities to the fixers.

BACKGROUND OF THE INVENTION

[0002] Curl and cockle of cellulose-based papers are persistent problems in inkjet printing with water-based inks. The problem stems from dimensional changes in the paper when it is wetted (especially when it is wetted on only one side) and then dried. In normal plain paper, dimensional stability is a function of the presence of cellulose fibers which are usually a couple of millimeters long. These bind together by fiber-to-fiber associations, which are dominated by inter-molecular H-bonds.

When these fiber-to-fiber H-bonds are disrupted or broken, changes in paper physical integrity are brought about. This breaking can be brought about by exposure to elevated temperatures, H-bonding solvents (including water) and/or moisture/humidity.

When aqueous fluid (ink/fixer) is applied to paper, it first accumulates in the paper's capillary spaces. Water and other hydrophilic components of the fluid wet the surfaces of the fibers. This water and/or organic cosolvent breaks the fiber-to-fiber H-bond associations and noticeably reduces the paper's dimensional integrity. Solvents that are particularly good at breaking the surface fiber-to-fiber interactions include water, dimethyl sulfoxide (DMSO), formamide, dimethyl formamide (DMF), ethylene glycol, glycerol, morpholine and pyridine.

[0003] With continued exposure of the aqueous-cosolvent fluid to the fibers in the paper, the water and hydrophilic solvents penetrate into the amorphous regions of the cellulose and cause the fibers to swell.

With wetting, the cellulose fiber-to-fiber associations (H-bonds) are disrupted by water and as the fibers swell with water, they increase in size, which relocates the original sites for fiber-to-fiber associations. As the fibers begin to dry from the outside inward, their fiber-to-fiber H-bonds tend to reestablish as surface moisture is lost. As the fibers continue drying out, they shrink from their swollen state, and with the surface fiber-to-fiber associations reestablished, stress/strain develops. This stress/strain is observed as curl across the page.

US5757408 discloses a recording sheet which, subsequent to being imaged with an aqueous ink, exhibit reduced curling.

This reduced curling is obtained by coating the paper substrate with a composition comprising 5-30 % wt. salt of an amino acid selected from β -alanine, glycine and γ -butyric acid. Said composition pertains to a fixer composition.

[0004] EP1524304 discloses an aqueous fixer fluid composition comprising a) 1-10 % wt. of an amino acid selected from β -alanine, 4-aminobutyric acid, 6-aminocaproic acid, L-aspartic acid, dimethylamino benzoic acid, and methyl-amino benzoic acid, b) a non-ionic water soluble solvent such as 1,2-hexanediol, c) a strong acid such as nitric acid, where the fluid has a pH of 4. The amino acids (a) have carboxylic acid functionality with pKa near 4.

SUMMARY OF THE INVENTION

[0005] One aspect of this disclosure includes an aqueous fixer fluid comprising at least one anti-curl cosolvent, wherein the at least one anti-curl cosolvent comprises at least one inner salt compound having a sp³- hybridized nitrogen cation and an associated anionic charge which is isolated on an oxygen anion bound directly to the sp³- hybridized nitrogen cation, wherein the at least one inner salt compound is an ammonium N-oxide moiety, and the fixer comprises a strong acid which results in an ammonium-N-oxide conjugate acid with an approximate pKa=4-5 and a buffering capacity in aqueous solution near pH=4-5, and wherein the aqueous fixer fluid comprises at least 10 weight percent of the at least one anti-curl cosolvent.

[0006] In addition, another aspect includes the use of an aqueous fixer fluid as defined herein to prevent paper curl in a printing process.

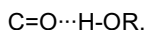
DETAILED DESCRIPTION OF EMBODIMENTS OF THE INVENTION

[0007] The present application relates to the problem of "curl" that results when cellulose-based media is inkjet printed with aqueous fluid. The present application describes employing "anti-curl" cosolvents that persist on the fiber surface and disrupt the fiber-to-fiber associations until the paper has dried and the state of swell of the cellulose has stabilized/equilibrated.

[0008] InkJet or other aqueous applications of fluid with either dyes, pigments or fixer on plain paper normally induces curl in the paper, depending upon the amount of fluid deposited on the paper, i.e. the amount per unit area of aqueous ink deposited on the paper. In a similar way, inkjet or other aqueous application of fixer on plain paper also induces curl. This apparently occurs because when the evaporating water of the aqueous fluid leaves the swollen cellulose fiber

surfaces of the substrate, shrinkage and stress-buildup of the fibers occurs, especially if the fibers are pinned closely together in their swollen configuration prior to complete reduction in the state of fiber swell.

[0009] An anti-curl agent can reduce the tendency of the imaged paper to curl as the aqueous deposit dries. An anti-curl agent appears to suppress the reformation of cellulose surface fiber-to-fiber H-bonding associations during the water drying process of a printed substrate. As set out in the present application, cosolvents have been found, for use as components of aqueous ink/aqueous fixer printing vehicles, that are effective anti-curl agents. All the cosolvents that are used in the present invention as anti-curl agents were selected to be disruptive to fiber-to-fiber H-bonds in cellulose-based media, i.e., to be H-bond accepting and/or H-bond donating solvents. Solvents as described above, such as water, have the capability to participate in an H-bond association in competition with the cellulose-cellulose H-bond associations. Such solvent powers of H-bond associations are quantitated in terms of H-bond accepting and H-bond donating. The H-bonds are asymmetric with a typical bond shown below with the C=O end being the acceptor and the RO-H being the donor:



For further explanation of H-bonding and its quantitation see: Progress in Physical Organic Chemistry, Vol. 13, R.W. Taft, ed., John Wiley, NY (1981), "An Examination of Linear Solvation Energy Relationships" M.J. Kamlet, J.L.M. Abboud, and R.W. Taft, page 485; M.J. Kamlet, J.L. M. Abboud, M.H. Abraham, and R.W. Taft, J. Org. Chem., Vol. 48, p2877-2887 (1983) "Linear Solvation Energy Relationships. 23. A Comprehensive Collection of the Solvatochromic Parameters, π^* , α , and β and Some Methods for Simplifying the Generalized Equation."

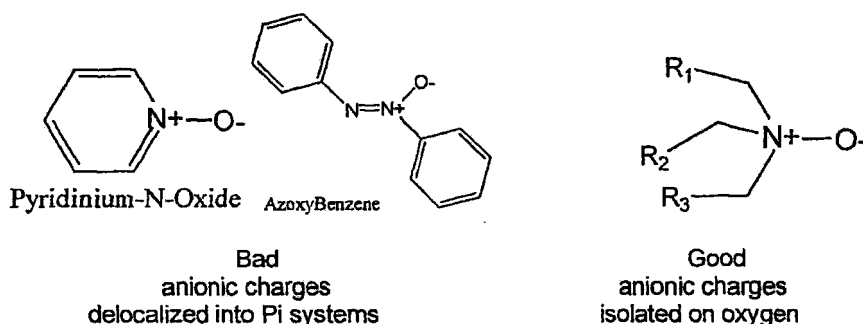
[0010] Non-limiting examples of known typical solvents/functional groups with strong H-bond accepting capabilities include: water, alcohols, ethers, amines, amides, carbonyl groups, sulfoxides, sulfones, phosphine oxides, etc. By the same token, non-limiting examples of typical solvents/functional groups with strong H-bond donating capabilities include: water, alcohols, protonated amines, primary and secondary amines, primary and secondary amides, etc.

[0011] In a preferred embodiment, the cosolvents used in the present invention also have low neat solvent vapor pressures, "neat solvent" being defined as an essentially pure, undiluted form of the solvent. Their anti-curl qualities appear to at least partially occur because their non-volatile character means they persist on the surface of the cellulose fibers during drying and continue to disrupt the fiber-to-fiber bonding until the excess water has evaporated and the fiber swell has stabilized.

[0012] The cosolvent compounds of the present invention all contain an inner salt. These inner salt-containing compounds are neutral compounds which have formal unit electrical charges of opposite sign. The inner-salt-containing compounds have relatively localized charges which are not part of an electronically conjugated system except in the case of conjugation between the oxygens within carboxylate groups. Specifically these compounds have a sp^3 -hybridized nitrogen cation, which cannot delocalize charge into a conjugated Pi electronic system, and an associated anionic charge isolated on an N-substituted oxygen anion (oxygen anion bound directly to the cationic nitrogen).

[0013] Ammonium-N-Oxides are shown below which fall inside and outside of the definition given above. The first one shown, pyridinium-N-oxide, has an sp^2 nitrogen cation and when tested, was found to be a poor anti-curl solvent. The second, azoxybenzene, is an "oxidized dye" which also has an sp^2 nitrogen cation. Azoxybenzene and pyridinium-N-oxide also have oxyanions that are part of a conjugated electronic system with expected anionic charge delocalization. Azoxybenzene, like the "Dimroth/Reichardt Betaine indicator dye", is not useful as an anti-curl solvent because it is not polar enough. These materials with sp^2 -hybridized nitrogen cations and oxyanions with delocalized charge are to be distinguished from the generalized structure of ammonium-N-oxide materials, (the third structure shown below), which has an sp^3 -hybridized nitrogen cation and an oxyanion with localized charge occurring in the "good" materials. The materials having this structure are useful and effective as anti-curl solvents.

[0014] In the case of these ammonium-N-Oxide materials, the desirable "good" example has a sp^3 -hybridized nitrogen cation that is not part of a conjugated Pi electronic system and the cationic charge is isolated on the Nitrogen. Similarly in the "good" example, the anionic charge on the oxygen is isolated to the oxygen, since the adjacent Nitrogen is fully saturated. In contrast, in the undesirable "bad" examples, the azoxybenzene and the pyridinium-N-oxide, the Nitrogens in them are sp^2 -hybridized and part of conjugated Pi electron systems, where the cationic charge is delocalized throughout the structure. Furthermore, the fact that the adjacent atom (Nitrogen) is sp^2 -hybridized allows the oxyanion to conjugate/delocalize into the extended Pi systems.



[0015] Non-limiting examples of inner salts in accordance with the invention which function well as anti-curl agents include ammonium-N-oxide moieties. Besides acting as anticurl agents, the oxyanions can act as conjugate bases for formation of weak acid buffers. These cosolvents in the presence of a strong acid, can also act as a buffer to provide substantial acid capacity in the presence of a strong acid, which is useful in fixers. This dual function can reduce the chemical loading in fixer formulations.

[0016] Non-limiting examples of ammonium-N-oxide moieties are: N-MethylMorpholine-N- Oxide (MMNO); N-Ethyl-Morpholine-N-Oxide (EMNO); N,N-DiMethylButylAmmonium-N-Oxide (DMBANO) and N,N,N-TriMethylAmmonium-N-Oxide (TMANO). Ammonium-N-oxide moieties have an oxygen anion and three alkyl groups attached to a cationic nitrogen, and each alkyl group independently comprises up to 6 carbons. Furthermore, each alkyl group is either straight-chained or branched, or, alternatively, multiple alkyl groups connected to the nitrogen may be combined together to form (a) 5 to 7- membered ring(s) comprising at least 4 carbons. Additionally, the alkyl groups may be substituted with or have attached to them groups such as water solubilizing moieties. For example, one or more of the carbons in a 5 or 6-membered ring can be substituted with an oxygen atom, such as the ether group in a morpholine ring or with a nitrogen atom. Other examples might include the attachment, rather than the substitution, of water solubilizing group(s) to alkyl group(s), that are in either straight-chained, branched or 5 or 6- membered ring groups. As a non-limiting example, the at least one water-solubilizing moiety might be a hydroxyl group, a carbonyl group, an amide group, a sulfone group a sulfoxide group, a polyethylene glycol moiety, or even an additional ammonium-N-oxide.

[0017] Of the ammonium-N-oxide moieties, Trimethylammonium-N-oxide (TMANO) is most preferred as an anti-curl agent. It has been found to have the best anti- curl qualities on a weight for weight basis of any of the ammonium-N-oxide cosolvents examined.

[0018] MMNO and the other ammonium-N-oxide cosolvents have also been found to be useful in their conjugate acid form (protonation of the N-Oxide anion, a weak base, in the presence of a strong acid) for buffering and increasing acid capacity in fixer vehicle. The "buffer" is in fact a mixture of the ammonium-N-oxide with its protonated form which is accessible by lowering the pH of the aqueous solution of the ammonium-N-oxide cosolvent with a strong acid (such as non-limiting examples: HCl, Nitric acid or MethaneSulfonic Acid) that, in the case of N-MethylMorpholine-N-Oxide, protonates the MMN^+O^- to form the acid MMN^+OH . Buffer capacity in the region of pH 4-5 is provided by ammonium-N-oxides which have conjugate acid pK_a 's (negative log acid dissociation constants) in the 4-5 region. This is the region that is best for maximizing safety and minimizing corrosion when the cosolvent is used in acidic fixers.

[0019] To maximize their anti-curl capabilities, any of the inner-salt-containing cosolvents or combination of the inner salt containing cosolvents of the present invention can be dissolved at levels of at least approximately 10% in an aqueous ink or fixer. Depending upon the formula weight of the inner salt cosolvent, 20-25 weight% is an optimum level of the cosolvent in an aqueous ink or fixer. The cosolvents can also be used at levels above approximately 25 weight %.

[0020] Any of the inner-salt-containing cosolvents of the present invention can also be dissolved in combination with other conventional non-ionic, water-soluble blendcosolvents in an aqueous solution. The total amount of the blend of the inner-salt-containing cosolvents and the non-ionic, water-soluble blend cosolvents comprises at least approximately 10 weight % of the aqueous ink or fixer vehicle. In a preferred embodiment, these blend cosolvents should preferably be, in addition to non-ionic and water soluble, H-bond accepting and/or H-bond donating. Such a combination of an inner-salt-containing cosolvent and blend cosolvent(s) can achieve anti-curl effects comparable to a similar concentration of the inner salts alone. In a more preferred embodiment, the non-ionic, water-soluble, H-bond accepting and/or H-bond donating, conventional cosolvent used should have a neat solvent vapor pressure of no more than 4×10^{-2} mm Hg (at 25°C). In a yet more preferred embodiment, the non-ionic, water-soluble, H-bond accepting and/or H-bond donating, conventional cosolvent used should have a neat solvent vapour pressure of no more than 1×10^{-4} mm Hg (at 25°C). In another preferred embodiment, the aqueous printing fluid comprises from 5 to 60% of the blended solution of the at least one inner-salt-containing cosolvent and the blend cosolvent(s) and the weight ratio in the blended solution of the inner-salt-containing cosolvent to blend cosolvent(s) ranges from 3:97 to 99:1.

[0021] Non-limiting examples of conventional cosolvents that can be blended with inner-salt-containing cosolvents are

many aqueous cosolvents conventionally used in inkjet printing fluids. Such conventional aqueous cosolvents might include, among others, at least one of the following: dipropylene glycol, 1,2-hexanediol, 2-pyrrolidinone, diethylene glycol, sulfolane, 1,5-pentanediol, tetraglyme, triethylene glycol, 1,6-hexanediol, glycerol, trimethylolpropane (EHPD), and tetraethylene glycol.

[0022] In a preferred embodiment, the inner-salt-containing cosolvent of the present invention or the blend of the inner-salt-containing cosolvent and the conventional cosolvent comprise from approximately 20 weight % to approximately 25 weight % of an aqueous ink or fixer vehicle. Inks and fixers formulated with these cosolvents interact well with components such as pigments, surfactants and colorants normally used in aqueous inkjet inks or fixers.

[0023] To further illustrate the present invention, the following examples are given. It is to be understood that these examples are provided for illustrative purposes and are not to be construed as limiting the scope of the present invention.

EXAMPLES

Example 1

[0024] This example compares anti-curl qualities of Ammonium-N-Oxides and Betaine with conventional cosolvents. Specifically, 7" X 9.5" images on 8.5"X11" sheets of Union Camp Great White paper (0.75" margins) were inkjet printed (10 6pL Drops/300 dpi) with aqueous printing fluid containing 15 weight %, 20 weight % and 25 weight % of one of the following cosolvents: DiEthyleneGlycol, TetraEthyleneGlycol, TriMethylolPropane (EHPD), 2-Pyrrolidinone, N-Methyl-Morpholine-N-Oxide, N-EthylMorpholine-N-Oxide, N,N-DiMethylButylAmine-N-Oxide, N,N,N-TriMethylAmine-N-Oxide, and Betaine. To facilitate thermal inkjet printing and a uniform image, the following additional components were included: 0.75% Tergitol 15S7, a surfactant; 0.05% EDTA, a metal sequestrant; and 0.05 % Acid Red 289, a trace magenta dye, to check image uniformity. These prints were allowed to dry unassisted at 23% relative humidity (RH) on a flat surface.

[0025] The curl of each sample was evaluated at 72 hours after printing. On Table 1 below, the curl of each sample was listed. With ambient drying, the curl of the worst edge rising above a flat surface was recorded in mm or, if the edge curled up and back toward the center of the paper, the curl was referred to as ">90". The description "Tube" was used when the curl resembled a scroll.

TABLE 1

	Curl (mm) at 72hr		
Water/No Cosolvent/Control			tube
	%Loading		
	15%	20%	25%
DiEthylene Glycol	tube	tube	tube
TetraEthylene Glycol	>90	45	25
EHPD	35	18	14
2-P	tube	tube	tube
MeMorpholine-N-Oxide	44	25	5
EtMorpholine-N-Oxide	40	20	5
DiMeBuAmine-N-Oxide	48	15	4
TriMeAmine-N-Oxide	35	7	1
Betaine (N,N,N-Trimethyl glycinate)	34	25	13

Example 2

[0026] This example compares anti-curl qualities of amino acids and Betaine with conventional cosolvents. Specifically, 7" X 9.5" images on 8.5"X11" sheets of Union Camp Great White paper were inkjet printed (10 6pL Drops/300 dpi) with aqueous printing fluid containing 15 weight %, 20 weight % and 25 weight % of one of the following cosolvents: 2-Pyrrolidinone, Glycerol, TetraEthyleneGlycol, 1,2-Hexanediol, Pyridine-N-Oxide, Betaine, α -Glycine, β -Alanine, and γ -AminoButyric Acid. To facilitate thermal inkjet printing and a uniform image, the following additional components were included: 0.75% Tergitol 15S7, a surfactant; 0.05% EDTA, a metal sequestrant; and 0.05 % Acid Red 289, a trace

magenta dye, to check deposit uniformity. Prints were allowed to dry unassisted at 35% RH on a flat surface.

[0027] The curl of each sample was evaluated at 72 hours after printing. On Table 2 below, the curl of each sample was listed in terms of mm curl, with curl in mm of worst edge rising above a flat surface with ambient drying. ">90" referred to the edge curling back toward the center of the paper. "Tube" referred to a scroll.

TABLE 2

	Curl (mm) at 72hr		
Water Control			tube
	% Loading		
CoSolvent	15%	20%	25%
2P	tube	Tube	tube
TetraEG	32	36	28
Glycerol	tube	Tube	tube
1,2-Hexanediol	tube	Tube	tube
Pyridine-N-Oxide	tube	Tube	tube
Betaine(N,N,N-Trimethyl Glycinate)	31	10	3
α -Glycine	41	31	Solubility Limit
β -Alanine	22	30	10
γ -Aminobutyric acid	32	26	2

Example 3

[0028] Curl data were collected for N-MethylMorpholine-N-Oxide and several other cosolvents at 15, 20, or 25% loading in aqueous inks also containing 0.75% Tergitol 15S7, 0.05% EDTA and 0.05% AR289 dye. These relatively pale inks (low colorant load) were printed on Union Camp Great White plain paper. These cosolvents were: Ethylene Glycol, Formamide, DiPropylene Glycol, 1,2-Hexanediol, 2-Pyrrolidinone, DiEthylene Glycol, Sulfolane, 1,5-Pentanediol, TriEthylene Glycol, 1,6-Hexanediol, Glycerol, TriMethylolPropane (EthylHydroxymethylPropaneDiol), TetraEthylene Glycol and N-MethylMorpholine-N-Oxide. Data were gathered at 1 hour (Table 3a) and 72 hours (Table 3b). Literature values of the neat solvent vapor pressures (calculated for 25°C) were quoted in mmHg. The curl of the imaged paper at 1 hour and 33% relative humidity (RH) and at 72 hrs and 43% RH was measured in millimeters (mm) of the height of the worst edge rising above a flat surface with non-assisted ambient drying. ">90" referred to the edge curling back toward the center of the paper. "Tube" referred to a scroll.

[0029] The data at 1 hour (Table 3a) below showed an improvement in 1 hr curl at vapor pressures below about 4×10^{-2} mmHg, which could be the high vapor pressure preferred limit for "mixture solvents". Examination of the data for 72 hours (Table 3b) showed additional curl with time for the more volatile cosolvents and indicates that the more preferred solvents will have vapor pressures below 1×10^{-4} mmHg.

TABLE 3a

	Curl (mm) at 1 hr			
	Neat Vapor Press (25°C) in mmHg			
Water Control	25			40
	% Loading			
CoSolvent		15%	20%	25%
	>4E-2			
Ethylene Glycol	9.2E-2	tube	tube	tube
Formamide	6.1E-2	tube	tube	tube

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(continued)

		<4E-2			
5	DiPropylene Glycol	3.2E-2	50	50	22
	1,2-Hexanediol	2.0E-2	tube	55	55
	2-Pyrrolidinone	1.3E-2	46	40	45
	DiEthyleneGlycol	5.6E-3	>90deg	53	40
10	Sulfolane	5.1E-3	40	>90deg	36
	1,5-Pentanediol	4.4E-3	40	49	29
	TriEthyleneGlycol	1.3E-3	41	31	45
15	1,6-Hexanediol	1.0E-3	34	35	22
	Glycerol	1.7E-4	37	40	19
		<1E-4			
	TriMethylolPropane	9.5E-5	43	25	22
20	TetraEthyleneGlycol	6.2E-7	22	24	15
	N-MethylMorpholine-N-Oxide	Internal salt	41	27	7

TABLE 3b

		Curl (mm) at 72 hr			
		Neat Vapor Press (25°C) in mmHg			
30	Water Control	25			tube
			% Loading		
	CoSolvent		15%	20%	25%
		>4E-2			
35	Ethylene Glycol	9.2E-2	tube	tube	tube
	Formamide	6.1E-2	tube	tube	tube
		<4E-2			
40	DiPropylene Glycol	3.2E-2	tube	tube	tube
	1,2-Hexanediol	2.0E-2	tube	tube	tube
	2-Pyrrolidinone	1.3E-2	>90deg	tube	Tube
	DiEthyleneGlycol	5.6E-3	tube	tube	tube
45	Sulfolane	5.1E-3	>90deg	tube	tube
	1,5-Pentanediol	4.4E-3	tube	tube	tube
	TriEthyleneGlycol	1.3E-4	45	49	45
50	1,6-Hexanediol	1.0E-4	48	53	52
	Glycerol	1.7E-4	tube	tube	tube
		<1E-4			
	TriMethylolPropane	9.5E-5	40	22	23
55	TetraEthyleneGlycol	6.2E-7	32	36	38
	N-MethylMorpholine-N-Oxide	Internal salt	41	24	6

Example 4

[0030] A fixer formulation commonly used in inkjet contained a cosolvent, an acidifying agent such as succinic acid (SA) and a cationic polymer mordant. The use of SA at pH 4 (near its first pKa) provided substantial acid capacity in the fixer, much like a true pH buffer. In this experiment MMNO was used as a replacement for the cosolvent in the fixer.

[0031] Preparation of the MMNO cosolvent based fixer formulations proceeded normally. Aqueous MMNO (15%) solutions as initially prepared were slightly alkaline. The MMNO was found to contain a trace of N-MethylMorpholine, "free amine" impurity, but MMNO was also found to be a weak base. The conjugate acid of MMNO, its protonated form, had a measured pK_a of about 4.2. Thus, addition of 4% SA, which normally lowers the pH of "simple/unbuffered" cosolvents to pHs between 2 and 3, only lowered the measured pH down to about 5.1. The buffering capacity of the weak base, MMNO, limited the pH drop to 5.1, while providing substantial "acid capacity". Methanesulfonic acid was also used to lower the pH of the MMNO formulations to 5.1 and further down to 4.0.

[0032] In addition to the buffering and anti-curl qualities that the MMNO imparts to the fixer formulations, these MMNO-containing fixer formulations had viscosity, decap and spit characteristics close to the original fixer formulations, using conventional solvents.

Reference Example 5

[0033] N-EthylMorpholine-N-Oxide was synthesized according to a procedure described in Johnson, U.S. Patent Number 1,144,048.

[0034] The oxidation of Ethyl Morpholine was done in a 500 ml 3-necked flask with a thermometer, condenser, equi-pressure addition funnel, magnetic stirrer, and temperature control, heating (heating mantle)/cooling (ice bath). Ethyl-Morpholine (172.6g) was placed into the flask and heated to 70°C.

[0035] Hydrogen peroxide (145.7 g of 35%) was placed in the addition funnel. Since the oxidation exotherms near 70°C, the dropwise addition of H₂O₂ was very carefully adjusted to maintain a temperature near 70°C. Occasional use of the ice bath was needed to keep the reaction from overheating during the approximately 2 hours required for the addition. The reaction mix was kept at 70°C (heating mantle) for 4 hours after completion of the addition. It was then allowed to stand overnight.

[0036] The reaction mix was then concentrated on a rotary evaporator (oil-less pump rated at 0.1 mm Hg and trap). Though the product did not drop out when acetone was added, the material finally solidified with time. The product was melted by heating it and triturated thoroughly with toluene to remove any free amine. The cooled solid was then pumped down in a vacuum oven. The solid product (N-EthylMorpholine-N-Oxide) was clean with about ½ mole water present.

Reference Example 6

[0037] The synthesis of N,N-DiMethylButylAmmonium-N-Oxide was virtually the same as the synthesis of N-Ethyl-Morpholine-N-Oxide set out in Example 5. N,N-DiMethylButylAmine (151.8 g) was placed in the flask and 145.7 g of 35% H₂O₂ was put in the addition funnel. After completion of the addition and 2 hours at 70°C, the mix was rotoevaporated using an oil-less pump. Water could only be removed very slowly. As a result, the product was collected as an aqueous solution that contains about 18% water.

Reference Example 7

[0038] The effect of blending conventional polar, water-soluble co-solvents of differing neat solvent vapor pressures with zwitterionic anti-curl co-solvents was tested in dye-based and pigment-based aqueous inks by printing a 7"X9.5" uniform area fill image on Union Camp Great White 8.5"X11" (plain paper) sheets at 10 (9pL) drops/300 dpi. 2.5% AcidRed-52(AR52) magenta (sulfurhodamine B) was the dye colorant for the dye based inks and 3.0% Cabotjet 200 black pigment (a carbon black pigment from Cabot, Inc., Billerica, Massachusetts) was the pigment colorant for the pigment-based inks. The formulations all contained 0.75% Tergitol 15S7 and 0.05% Na₂EDTA. Two anti-curl cosolvents, 4-AminoButyric Acid (AmBuAcid) and N-MethylMorpholine-N-Oxide (MMNO) were selected as cosolvents (at 15% loading) for the dye-based and pigment-based inks respectively. At 15% loadings of the anti-curl agent, the curl of samples printed with this level of anti-curl agent was not fully suppressed. The contributions of inks containing 15% anti-curl solvent + additional cosolvents were compared to the 15% anti-curl controls. The calculated neat solvent vapor pressures of the additional cosolvents were also tabulated based upon literature data and associated formula and the selected temperature of 25°C. (See Chemical Properties Handbook Yaws, C.L. McGraw-Hill 1999 <http://www.knovvel.com/knovet2/Toc.jsp?SpaceID=10093&BookID=49>)

[0039] The dye-based ink comparisons using AmBuAcid anti-curl agent were collected in Table 7a. It was apparent that the curl can actually be made worse by blending a volatile cosolvent such as Ethylene Glycol (EG) with the anti-

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curl cosolvent. Generally, with 15% of the anticurl cosolvent, AmBuAcid, the curl position improved as the volatility of the blended 10% polar/H-bonding cosolvent is reduced. For cosolvents of modestly low volatility, such as 10% TriEthylene Glycol, compared to the control, the 1 hr curl was attenuated in a somewhat additive fashion (15% AmBuAcid + 10% TEG) and the small adverse impact of the solvent volatility became noticeable at 72hrs. For the essentially non-volatile cosolvents, the curl increases with extended dry time are barely noticeable (few mm).

[0040] In measuring the curl, the curl in mm was the height of the worst edge rising above a flat surface with ambient drying. ">90" referred to the edge curling back toward the center of the paper. "Tube" referred to a scroll.

Table 7a

Dye Based Inks with 15% β-AminoButyric Acid + 10% Additional Cosolvent			
		Curl in mm	
10% Additional CoSolvent	Neat Vapor Pressure (25°C) in mmHg	1 Hr	72Hrs
None		22	27
10% Ethylene Glycol	9.2E-2	28	41
10% 2-Pyrrolidinone	2.0E-2	14	>90
10% TriEthyleneGlycol	1.3E-3	9	27
10% TriEMethylolPropane	9.5E-5	6	10
10% TetraEthyleneGlycol	6.2E-7	5	13
10% 4-AminoButyric Acid	Internal Salt	5	7
Dye Based Inks with No Inner Salt, Just 10% "Additional" Cosolvent			
		Curl in mm	
10% Additional CoSolvent	Neat Vapor Pressure (25°C) in mmHg	1Hr	72Hrs
10% Water/Control	2.4E+1	36	39
10% Ethylene Glycol	9.2E-2	>90	>90
10% 2-Pyrrolidinone	2.0E-2	37	>90
10% TriEthyleneGlycol	1.3E-3	26	33
10% TriMethylolPropane	9.5E-5	18	18
10% TetraEthyleneGlycol	6.2E-7	>90	40
10% 4-AminoButyric Acid	Internal Salt	31	33

[0041] The corresponding CABOJET 200 pigment based ink comparisons using 15% N-MethylMorpholine-N-Oxide (MMNO) anti-curl agent were collected in Table 7b. As with the dye based ink series, the incorporation of a modestly low volatility cosolvent, TriEthylene Glycol, yielded increasing curl with extended dry time from 7mm at 1hr to 18mm at 72 hrs. There appeared to be some slow evaporation of cosolvent and migration out of the partially swollen cellulose, that manifests itself as delayed curl.

Table 7b

Pigmented Inks with 15% N-MethylMorpholine-N-Oxide + 10% Cosolvent			
		Curl in mm	
10 % Additional CoSolvent	Neat Vapor Pressure (25°C) in mm Hg	1 Hr.	72 Hr.
None		24	24
10% Ethylene Glycol	9.2E-2	40	>90
10% 2-Pyrrolidinone	2.0E-2	11	>90
10% TriEthyleneGlycol	1.3E-3	7	18

(continued)

Pigmented Inks with 15% N-MethylMorpholine-N-Oxide + 10% Cosolvent			
		Curl in mm	
10 % Additional CoSolvent	Neat Vapor Pressure (25°C) in mm Hg	1 Hr.	72 Hr.
10% TriMethylolPropane	9.5E-5	10	10
10% TetraEthyleneGlycol	6.2E-7	10	13
10% MMNO	Internal Salt	4	5
Pigmented Inks with No Inner Salt, Just 10% "Additional" Cosolvent			
		Curl in mm	
10% Additional CoSolvent	Neat Vapor Pressure (25°C) in mm Hg	1 Hr.	72 Hr.
10% Water	24	>90	>90
10% Ethylene Glycol	9.2E-2	tube	tube
10% 2-Pyrrolidinone	2.0E-2	>90	>90
10% TriEthyleneGlycol	1.3E-3	>90	>90
10% TriMethylolPropane	9.5E-5	26	30
10% TetraEthyleneGlycol	6.2E-7	40	>90

[0042] The AR-52 dye based inks with 15% 4-AminoButyric Acid incorporated as anti-curl agent were challenged with varying concentrations of the "relatively volatile" solvent Ethylene Glycol to determine the level at which its presence causes adverse changes in the paper curl. This data was collected in Table 7c. Examination of the 72hr data in Table 7c clearly showed a deleterious effect even at Ethylene Glycol concentrations as low as 3%.

Table 7c

Curl With 15% 4-AminoButyric Acid + Varying Levels of Ethylene Glycol					
		Curl in mm			
Additional CoSolvent	Neat Vapor Pressure (25°C) in mm Hg	1 Hr.	24 Hrs.	48 Hrs.	72 Hrs.
0% Ethylene Glycol		22	23	23	27
3% Ethylene Glycol	9.2E-2	29	34	34	36
6% Ethylene Glycol	9.2E-2	34	40	42	44
10% Ethylene Glycol	9.2E-2	28	37	40	41
15% Ethylene Glycol	9.2E-2	>90	>90	>90	tube
10% 4-AminoButyric Acid	Internal Salt	5	5	6	7

Claims

1. An aqueous fixer fluid comprising at least one anti-curl cosolvent, wherein the at least one anti-curl cosolvent comprises at least one inner salt compound having a sp^3 -hybridized nitrogen cation and an associated anionic charge which is isolated on an oxygen anion bound directly to the sp^3 -hybridized nitrogen cation, wherein the at least one inner salt compound is an ammonium N-oxide moiety, and the fixer comprises a strong acid which results in an ammonium-N-oxide conjugate acid with an approximate $pK_a=4-5$ and a buffering capacity in aqueous solution near $pH=4-5$, and wherein the aqueous fixer fluid comprises at least 10 weight percent of the at least one anti-curl cosolvent.
2. The aqueous fixer fluid of claim 1, wherein the at least one anti-curl cosolvent forms a blended solution with at least

one non-ionic, water-soluble blend cosolvent.

3. The aqueous fixer fluid of claim 2, wherein the at least one blend cosolvent has a neat solvent vapor pressure of no more than 4×10^{-2} mm Hg (5.3 Pa) and is capable of strong H-bonding interactions for disrupting fiber-to-fiber H-bonds in cellulose-based paper.
4. The aqueous fixer fluid of claim 3, wherein the at least one blend cosolvent has a neat solvent vapor pressure of no more than 1×10^{-4} mm Hg (1.3×10^{-2} Pa).
5. The aqueous fixer fluid of claim 2, wherein the aqueous fixer fluid comprises from 5 to 60% of the blended solution of the at least one anti-curl cosolvent and the at least one blend cosolvent and the weight ratio of the inner salt anti-curl cosolvent(s) to blend cosolvent(s) in the blended solution ranges from 3:97 to 99:1.
6. The aqueous fixer fluid of claim 5, wherein the at least one inner salt compound comprises an ammonium N-oxide moiety with three alkyl groups attached to the nitrogen, and wherein each alkyl group independently comprises up to 6 carbons, each alkyl group being either a straight-chain, a branched chain, or a 5 to 7-membered ring comprising at least 4 carbons.
7. The aqueous fixer fluid of claim 6, wherein the 5 to 7-membered ring comprises at least one hetero atom selected from the group consisting of oxygen and nitrogen.
8. The aqueous fixer fluid of claim 7, wherein the 5 to 7-membered ring is a morpholine ring.
9. The aqueous fixer fluid of claim 6, wherein at least one of the three alkyl groups comprises at least one water-solubilizing substituent.
10. The aqueous fixer fluid of claim 9, wherein the at least one water-solubilizing substituent is selected from the group consisting of a hydroxyl group, a carbonyl group, an amide group, a sulfone group, a sulfoxide group, a polyethylene glycol moiety and an ammonium-N-oxide moiety.
11. The aqueous fixer fluid of claim 1, wherein the at least one inner salt compound is selected from the group consisting of N-MethylMorpholine-N-Oxide (MMNO); N-EthylMorpholine-N-Oxide (EMNO); N,N-DiMethylButylAmmonium-N-Oxide (DMBANO) and N,N,N-TriMethylAmmonium-N-Oxide (TMANO).
12. The use of an aqueous fixer fluid as claimed in any preceding claim to prevent paper curl in a printing process.

Patentansprüche

1. Wässriges Fixiermittelfluid, das wenigstens ein Anti-Kräuseln-Co-Lösungsmittel umfasst, wobei das wenigstens eine Anti-Kräuseln-Co-Lösungsmittel wenigstens eine innere Salzverbindung mit einem sp^3 -hybridisierten Stickstoffkation und einer verknüpften anionischen Ladung umfasst, die auf einem Sauerstoffanion, das direkt an das sp^3 -hybridisierte Stickstoffkation gebunden ist, isoliert ist, wobei die wenigstens eine innere Salzverbindung ein Ammonium-N-Oxid-Anteil ist und das Fixiermittel eine starke Säure umfasst, was zu einer konjugierten Ammonium-N-Oxid-Säure mit einer ungefähren $pK_a=4-5$ und einer Pufferfähigkeit in einer wässrigen Lösung nahe einem $pH=4-5$ führt, und wobei das wässrige Fixiermittelfluid wenigstens 10 Gewichtsprozent des wenigstens einen Anti-Kräuseln-Co-Lösungsmittels umfasst.
2. Wässriges Fixiermittelfluid nach Anspruch 1, wobei das wenigstens eine Anti-Kräuseln-Co-Lösungsmittel eine gemischte Lösung mit wenigstens einer nicht ionischen, wasserlöslichen Co-Lösungsmittelmischung ausbildet.
3. Wässriges Fixiermittelfluid nach Anspruch 2, wobei die wenigstens eine Co-Lösungsmittelmischung einen reinen Lösungsmitteldampfdruck von nicht mehr als 4×10^{-2} mm Hg (5,3 Pa) aufweist und zu starken H-bindenden Interaktionen zum Unterbrechen von Faser-zu-Faser-H-Bindungen in Papier auf Zellulosebasis fähig ist.
4. Wässriges Fixiermittelfluid nach Anspruch 3, wobei die wenigstens eine Co-Lösungsmittelmischung einen reinen Lösungsmitteldampfdruck von nicht mehr als 1×10^{-4} mm Hg ($1,3 \times 10^{-2}$ Pa) aufweist.

5. Wässriges Fixiermittelfluid nach Anspruch 2, wobei das wässrige Fixiermittelfluid von 5 bis 60 % der gemischten Lösung des wenigstens einen Anti-Kräuseln-Co-Lösungsmittels und der wenigstens einen Co-Lösungsmittelmischung umfasst und das Gewichtsverhältnis des/der inneren Salz-Anti-Kräuseln-Co-Lösungsmittel(s) zu der/den Co-Lösungsmittelmischung(en) in der gemischten Lösung von 3:97 bis 99:1 reicht.
6. Wässriges Fixiermittelfluid nach Anspruch 5, wobei die wenigstens eine innere Salzverbindung einen Ammonium-N-Oxid-Anteil mit drei Alkylgruppen umfasst, die an dem Stickstoff angelagert sind, und wobei jede Alkylgruppe unabhängig bis zu 6 Kohlenstoffatome umfasst, jede Alkylgruppe entweder eine gerade Kette, eine verzweigte Kette oder ein 5 bis 7-gliedriger Ring, wenigstens 4 Kohlenstoffatome umfassend, ist.
7. Wässriges Fixiermittelfluid nach Anspruch 6, wobei der 5 bis 7-gliedrige Ring wenigstens ein Heteroatom umfasst, ausgewählt aus der Gruppe bestehend aus Sauerstoff und Stickstoff.
8. Wässriges Fixiermittelfluid nach Anspruch 7, wobei der 5 bis 7-gliedrige Ring ein Morpholin-Ring ist.
9. Wässriges Fixiermittelfluid nach Anspruch 6, wobei wenigstens eine der drei Alkylgruppen wenigstens einen Wasser löslich machenden Substituenten umfasst.
10. Wässriges Fixiermittelfluid nach Anspruch 9, wobei der wenigstens eine Wasser löslich machende Substituent ausgewählt ist aus der Gruppe bestehend aus einer Hydroxylgruppe, einer Carbonylgruppe, einer Amidgruppe, einer Sulfongruppe, einer Sulfoxidgruppe, einem Polyethylenglycolanteil und einem Ammonium-N-Oxid-Anteil.
11. Wässriges Fixiermittelfluid nach Anspruch 1, wobei die wenigstens eine innere Salzverbindung ausgewählt ist aus der Gruppe bestehend aus N-Methylmorpholin-N-Oxid (MMNO); N-Ethylmorpholin-N-Oxid (EMNO); N,N-Dimethylbutylammonium-N-Oxid (DMBANO) und N,N,N-Trimethylammonium-N-Oxid (TMANO).
12. Verwendung eines wässrigen Fixiermittelfluids nach einem der vorhergehenden Ansprüche, um ein Papier-Kräuseln in einem Druckvorgang zu verhindern.

Revendications

1. Liquide fixateur aqueux comprenant au moins un co-solvant anti-incurvation, dans lequel l'au moins un co-solvant anti-incurvation comprend au moins un composé de sel interne ayant un cation azote hybridé sp^3 et une charge anionique associée qui est isolée sur un anion oxygène lié directement au cation azote hybridé sp^3 , dans lequel l'au moins un composé de sel interne est un fragment N-oxyde d'amine, et le fixateur comprend un acide fort qui donne un acide conjugué N-oxyde d'amine avec un pK_a approximatif = 4 à 5 et une capacité tampon dans une solution aqueuse proche de $pH = 4$ à 5, et dans lequel le liquide fixateur aqueux comprend au moins 10 % en poids de l'au moins un co-solvant anti-incurvation.
2. Liquide fixateur aqueux selon la revendication 1, dans lequel l'au moins un co-solvant anti-incurvation forme une solution mélangée avec au moins un co-solvant de mélange non ionique hydrosoluble.
3. Liquide fixateur aqueux selon la revendication 2, dans lequel l'au moins un co-solvant de mélange a une pression de vapeur de solvant pure non supérieure à 4×10^{-2} mm Hg (5,3 Pa) et est capable de fortes interactions de liaison H pour désorganiser les liaisons H fibre-à-fibre dans le papier à base de cellulose.
4. Liquide fixateur aqueux selon la revendication 3, dans lequel l'au moins un co-solvant de mélange a une pression de vapeur de solvant pure non supérieure à 1×10^{-4} mm Hg ($1,3 \times 10^{-2}$ Pa).
5. Liquide fixateur aqueux selon la revendication 2, dans lequel le liquide fixateur aqueux comprend de 5 à 60 % de la solution mélangée de l'au moins un co-solvant anti-incurvation et de l'au moins un co-solvant de mélange et le rapport en poids du ou des co-solvant(s) contenant un sel interne au(x) co-solvant(s) de mélange dans la solution mélangée se situe dans la plage allant de 3:97 à 99:1.
6. Liquide fixateur aqueux selon la revendication 5, dans lequel l'au moins un composé de sel interne comprend un fragment N-oxyde d'amine avec trois groupes alkyles liés à l'azote, et dans lequel chaque groupe alkyle comprend indépendamment jusqu'à 6 atomes de carbone, chaque groupe alkyle étant soit une chaîne linéaire, soit une chaîne

ramifiée, soit un cycle de 5 à 7 chaînons comprenant au moins 4 atomes de carbone.

7. Liquide fixateur aqueux selon la revendication 6, dans lequel le cycle de 5 à 7 chaînons comprend au moins un hétéroatome choisi dans le groupe constitué par l'oxygène et l'azote.

8. Liquide fixateur aqueux selon la revendication 7, dans lequel le cycle de 5 à 7 chaînons est un cycle morpholine.

9. Liquide fixateur aqueux selon la revendication 6, dans lequel au moins l'un des trois groupes alkyles comprend au moins un substituant solubilisant dans l'eau.

10. Liquide fixateur aqueux selon la revendication 9, dans lequel l'au moins un substituant solubilisant dans l'eau est choisi dans le groupe constitué par un groupe hydroxyle, un groupe carbonyle, un groupe amide, un groupe sulfone, un groupe sulfoxyde, un fragment polyéthylène glycol et un fragment N-oxyde d'amine.

11. Liquide fixateur aqueux selon la revendication 1, dans lequel l'au moins un composé de sel interne est choisi dans le groupe constitué par l'oxyde de N-méthylmorpholine (MMNO) ; l'oxyde de N-éthylmorpholine (EMNO) ; l'oxyde de N,N-diméthylbutylamine (DMBANO) et l'oxyde de N,N,N-triméthylamine (TMANO).

12. Utilisation d'un liquide fixateur aqueux selon l'une quelconque des revendications précédentes pour empêcher l'incurvation du papier dans un processus d'impression.

REFERENCES CITED IN THE DESCRIPTION

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